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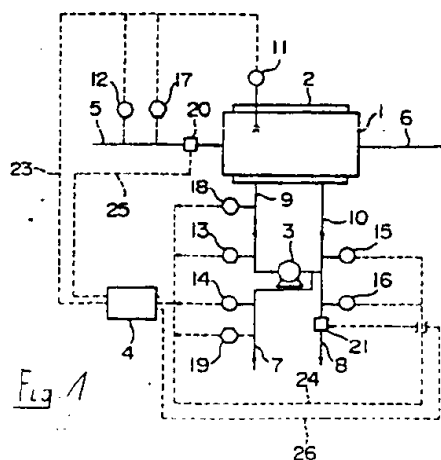
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(54) Process for producing a vinyl polymer.

(57) In a process for producing an intended polymer by polymerizing at least one vinyl compound, an improvement which comprises calculating the concentration of the unreacted vinyl compound in the reactor, the composition of the polymer formed, and the amount thereof from the amount of heat calculated from the heat balance of the reactor, the amounts of raw materials fed to the reactor, the reactivity ratio of the vinyl compounds and the heat of polymerization per unit amount of each of the vinyl compounds, and regulating the amounts of raw materials to be fed into the reactor in such a way that the values thus calculated agree respectively with the predetermined target values of the polymer composition and the amount of polymer formed.



PROCESS FOR PRODUCING A VINYL POLYMER

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1 This invention relates to a process for producing
a vinyl polymer, and more particularly, to a process for
producing an intended vinyl polymer stably.

 In the production of a vinyl polymer, it is very
5 important for obtaining the intended polymer stably to
maintain constant various factors of polymerization condi-
tions as polymerization temperature, polymerization pres-
sure, residence time, as well as concentrations of
catalysts and feed materials in the reaction system.
10 However, although some of these polymerization factors, for
example the reaction temperature, can be detected and
controlled in a relatively simple manner, there are some
other factors, for example the concentration of feed
material, which vary in the reaction system and are
15 difficult to detect and control directly. Especially when
two or more vinyl compounds are copolymerized, even
if the polymerization temperature, polymerization pres-
sure, and the feeding ratio of the vinyl compounds are
kept constant, the concentrations of the unreacted monomers
20 in the reaction system vary depending on the polymerization
activity of the catalyst and the concentration thereof,
and hence the composition of the resulting polymer
fluctuates.

 For controlling the polymerization of vinyl
25 compounds, there has been known a method previously

1 wherein the polymerization temperature, pressure and the
like are kept constant, some suitable component in the
reaction system is withdrawn and analyzed, and the
amounts of materials to be fed in the system are regulated
5 based on the result of the analysis. But this method is
not desirable because the sampling of a specimen containing
a polymer and the analysis of the components contained
therein are very troublesome, and a time lag due to
the time required for analysis is involved. When the
10 polymerization system has a gas phase free from the polymer,
the composition of the gas phase can be known by analysis
relatively easily. But, on converting the composition of
the gas phase into that of the system in which the polymeri-
zation is actually taking place, the accurate value of the
15 latter is often difficult to obtain owing to the limitation
due to mass transfer. Accordingly, the above method is
not satisfactory for controlling the polymerization
process.

There is also known a method which controls
20 the polymerization process by measuring the heat
evolved in polymerization. But this known method (Japanese
Patent Publication No. 13047/60) only regulates the catalyst
feeding rate so as to keep the rate of polymer formation
constant, which latter rate is learnt from the knowledge
25 of the heat evolved in the polymerization. This method
can not give stably a copolymer having a fixed composition
in copolymerization.

The present inventors have made extensive

1 studies on the control of such polymerization of vinyl
compounds, and as a result, established a technique to
control the polymerization easily and accurately, and
attained to this invention.

5 An object of this invention is to provide a
process for producing a vinyl polymer by which an
intended polymer can be produced stably.

Other objects and advantages of this invention
will become apparent from the following descriptions.

10 Thus, according to this invention, there is
provided, in a process for producing an intended polymer by
polymerizing at least one vinyl compound, an improvement
which comprises calculating the concentration of the unre-
acted vinyl compound in the reactor, the composition of the
15 polymer formed, and the amount thereof from the amount of
reaction heat calculated from the heat balance of the re-
actor, the amounts of raw materials fed to the reactor, the
reactivity ratio of the vinyl compounds and the heat of
polymerization per unit amount of each of the vinyl
20 compounds, and regulating the amounts of raw materials
to be fed into the reactor in such a way that the values
thus calculated agree respectively with the predeter-
mined target values of the polymer composition and the
amount of polymer formed.

25 This invention will be described further in
detail below.

The polymerization of a vinyl compound using
a catalyst is an exothermic reaction, and the value of
evolved heat is characteristic to the individual vinyl

1 compound to be polymerized. Further, the relation
between the concentrations of the vinyl compounds in the
reaction system and the composition of the polymer obtained
is in general determined unequivocally so long as the
5 kind of the catalyst system and the polymerization condi-
tions are kept constant. This invention provides, taking
advantage of these situations, a process for producing a
vinyl polymer, wherein the concentration of the unreacted
vinyl compound in the reactor is determined from the heat
10 and material balance of the reactor, and the amount of
raw materials to be fed is regulated so as to keep the
above concentration of the unreacted vinyl compound
constant, thus to produce the intended polymer stably.

The accompanying drawing is a flow sheet illust-
15 rating an example of the embodiment of this invention. The
process of this invention is explained below with reference
to the drawing.

The polymerization is carried out in a reactor
1 provided with a heat-removing jacket 2. The raw materials
20 are fed into the reactor through a line (or plural lines)
5, and the reaction mixture is withdrawn therefrom through
a line 6 in an amount corresponding to the fed amount.
A fixed amount of cooling water is circulated through
the reactor jacket 2 via lines 9 and 10 by means of
25 a circulating pump 3. Cooling water of higher temperature
is withdrawn through a line 8 out of the system, and cooling
water of lower temperature in an amount corresponding to
that of the former is introduced into the system through

1 a line 7.

The temperatures of the reactor and each lines are measured with thermometers 11 to 16. The flow rates in the lines are measured with flow meters 17 to 19 as required.

5 These measured values of temperatures and flow rates are input into a computer 4 via lines for instrumentation 23 and 24. The calculated results are compared with set values, and output signals to match the set values are transmitted through lines 25 and 26 to a flow controller 20.
10 for raw materials and a flow controller 21 for the discharged water in line 8, to control the respective flow rates. The line 5 is shown here to represent the feed lines for various raw materials as the catalyst, diluent for polymerization and vinyl compounds collectively. The
15 temperature measurements, flow rate measurements and flow controls can be made for each of the feed lines as required. The measurement of flow rate in each line can be made by any suitable methods including a method to use a continuous measuring instrument such as an orifice-type
20 flowmeter or a rotameter, one to detect the level change of a metering pipe, or one to charge batchwise by use of a metering pipe. Similarly, various methods can be used to control the flow rate in each line. The control may be effected by such methods as one to use a conventional
25 flow control valve, one to charge batchwise by use of a metering pipe or, when the raw materials are introduced by means of a reciprocating pump, a method to regulate the speed of reciprocating motion.

1 In the computer 4, by using the data of the
temperature and flow rate input from each line, physical
characteristics of each raw material input beforehand (for
example enthalpy), heat of polymerization of each vinyl
5 compound, and reactivity ratio of each vinyl compound,
the heat balance and the material balance are determined,
the amount and composition of the polymer formed are calcu-
lated, and the values obtained are compared with the set
values to output signals for the control instruments 20
10 and 21.

 The heat removed from the polymerization
reactor can be calculated from the flow rates of cooling
waters through lines 7 and 8 and the difference in
temperatures thereof. But the temperature difference (ΔT)
15 between the inner temperature of the reactor 1 and that of
jacket 2 can also be used as a parameter. Namely, ΔT
assumes a value corresponding to the amount of evolved
heat when the overall heat transfer coefficient at the
reactor wall is constant. Since an abrupt change in
20 the evolving heat shows itself first as a change in ΔT ,
 ΔT can be used as the parameter for controlling reaction
for a short-time.

 If necessary, for the calculation of the heat
balance, other physical quantities such as pressure,
25 though not shown in the drawing, can be measured at
each line or apparatus, and input to the computer 4.
The method for determining the heat and material balance
and the method for calculating the amount and composition

1 of polymer formed are described concretely below.

Total amount of removed heat is calculated from the flow rates and temperature difference of cooling waters through lines 7 and 8. From this value, are subtracted
5 the amounts of such heat other than that due to polymerization as the sensible heats of fed materials, heat generated by stirring, and heat released outside, to give the total amount of heat evolved by polymerization per unit time. This heat amount is expressed as Q .

10 It is assumed that, of the raw materials fed into the reaction system, i kinds of vinyl compounds (hereinafter referred to as "monomers") participate in the reaction. Of these i monomers, the fed amount, unreacted amount, and polymerized amount per unit
15 time are put as follows respectively:

$M_i(1)$: Fed amount of the i -th monomer,

$M_i(2)$: Unreacted amount of the i -th monomer,

$M_i(3)$: Polymerized amount of the i -th monomer.

Further, the heat of polymerization of the
20 i -th monomer per unit amount is put as h_i .

Firstly, from the heat balance holds the following equation:

$$Q = h_1 M_1(3) + h_2 M_2(3) + h_3 M_3(3) + \dots + h_i M_i(3).$$

From the material balance follows the i equations below:

- 8 -

$$M_1(1) = M_1(2) + M_1(3)$$

$$M_2(1) = M_2(2) + M_2(3)$$

$$\vdots$$

$$M_i(1) = M_i(2) + M_i(3)$$

- 1 Then, when the polymer composition is expressed taking the amount of the first monomer as 1 and that of the i -th monomer as a_i , the following $i-1$ equations hold:

$$M_2(3) = a_2 M_1(3)$$

$$M_3(3) = a_3 M_1(3)$$

$$\vdots$$

$$M_i(3) = a_i M_1(3)$$

- 5 Similarly, when the monomer composition is expressed taking the amount of the first monomer as 1 and that of the i -th monomer as b_i , the following $i-1$ equations hold;

$$M_2(2) = b_2 M_1(2)$$

$$M_3(2) = b_3 M_1(2)$$

$$\vdots$$

$$M_i(2) = b_i M_1(2)$$

1 Further, when the relationship between the monomer composition and the polymer composition is determined beforehand by a separate experiment, also $i-1$ equations hold as follows:

$$\begin{aligned} a_2 &= f_2(b_2, b_3, \dots, b_i) \\ a_3 &= f_3(b_2, b_3, \dots, b_i) \\ &\vdots \\ a_i &= f_i(b_2, b_3, \dots, b_i) \end{aligned}$$

5 From the foregoing, there hold $1+i+(i-1)+(i-1)+(i-1) = 4i-2$ equations. In these equations, $M_1(1), M_2(1), \dots, M_i(1)$ are known, being already measured at line 5, and h_1, h_2, \dots, h_i are data representing physical properties, namely known values.

10 Accordingly, the number of unknowns are as follows:

$M_1(2), M_2(2) \dots, M_i(2)$	i
$M_1(3), M_2(3) \dots, M_i(3)$	i
a_2, a_3, \dots, a_i	$i-1$
b_2, b_3, \dots, b_i	$i-1;$

namely $4i-2$ in sum. The amount of unreacted monomer and the polymer composition are obtained by solving these equations. However, the relationship between the monomer
15 composition and the polymer composition can also be

1 represented not in a mathematical expression as above, but
in a graph. In this case, the intended values can be
obtained by determining a monomer composition from the
graph assuming a value for polymer composition, sub-
5 stituting these values for corresponding letters in the
material balance and heat balance equations, and repeating
the assumption of the polymer composition and the calcu-
lation of material and heat balance until no inconsistency
can be observed. Such repetitive calculations can be
10 conducted advantageously by the use of the computer 4.

In any case, by measuring the heat evolved in
polymerization as well as the amount of fed monomer, the
amount of unreacted monomer, the amount of polymer, and the
compositions thereof can be calculated based on the
15 knowledge of the monomer reactivity ratio (the relation-
ship between the composition of unreacted monomer and
that of polymer formed). Naturally, this invention can
also be applied to the case wherein $i = 1$, namely to the
homopolymerization.

20 The amount and composition of the polymer
thus obtained are compared with the objective
values for production, and the amounts of the catalyst
and the monomer to be fed are regulated accordingly to
achieve the objective values.

25 Further, it is possible to calculate the heat
which will be evolved in polymerization beforehand from
the objective values for production and the amount of a
suitable combination of monomers to be fed, and to regulate

1 the amount of catalyst to be fed such that actually
evolved heat agrees with that calculated above. But when
some deviation from the calculated value of evolving heat
occurs, it is very important to know what composition of
5 polymer is being obtained. Also for attaining to the
objective composition most efficiently, it is necessary
to know what composition of polymer is being obtained.

Following the procedures mentioned above, the
intended polymer can be produced stably.

10 This invention can be applied to various polymeri-
zations of vinyl compounds including radical, cationic,
and anionic polymerization. The radical polymerization
is effected by the action of peroxides, azo compounds,
redox initiators and the like, and is conducted in the
15 form of bulk, solution, emulsion, or suspension poly-
merization. Representative examples of compounds subjected
to radical polymerization include ethylene, styrene,
vinyl chloride, vinylidene chloride, acrylonitrile, vinyl
acetate, and methyl methacrylate.

20 The cationic polymerization is initiated by
protonic acids such as hydrogen chloride and sulfuric acid,
metal halides such as BF_3 , AlCl_3 and TiCl_4 , metal alkyls
such as $\text{AlC}_2\text{H}_5\text{Cl}_2$ and $\text{Al}(\text{C}_2\text{H}_5)_3$, and acidic oxides such
as CrO_3 , MoO_3 , and $\text{SiO}_2\text{-Al}_2\text{O}_3$. The compounds subjected to
25 cationic polymerization include, for example, styrene,
isobutene and alkyl vinyl ether.

The anionic polymerization uses as catalyst
alkali metals, alkali metal complexes, organometallic

1 compounds and the like. Examples of compounds subjected
to anionic polymerization include styrene, butadiene,
methyl methacrylate, acrylonitrile and vinylidene
cyanide.

5 This invention can be applied further to the
polymerization catalyzed by a combination of a transition
metal compound and an organometallic compound, the so-
called Ziegler-Natta catalyst. The polymerization by
Ziegler-Natta catalyst uses a compound (halides, oxyhalides,
10 esters etc.) of a transition metal of the groups IV to VIII
and an organometallic compound of a metal of the groups I
to III of the periodic table as the catalyst. There are
also used catalyst systems wherein the transition metal
compound is supported on a magnesium compound, silica,
15 alumina and the like to enhance the catalytic activity.

The monomers which can be used in this invention
include α -olefins such as ethylene, propylene, butene-1,
pentene-1, hexene-1, and 4-methylpentene-1 as well as
dienes such as butadiene and isoprene, and polar vinyl
20 compounds such as methyl methacrylate, vinyl ether and
vinyl chloride. The polymerization can be carried out
by any of the methods of bulk, solution, slurry, and
gas-phase polymerization.

The polymerization conditions for the process
25 of this invention are not restricted specifically, and
may be suitably selected in accordance with the method of
polymerization adopted. This invention may be practiced
in a single reactor, but it may also be applied to plural

1 reactors connected in series, namely to so-called series
polymerization. In the series polymerization, when poly-
mers of different compositions are formed in each of the
reactors, it is impossible to know individually the
5 respective polymer composition being formed in each of the
reactors by analyzing merely the polymer obtained in the
latter step of polymerization, and hence it is difficult
to control individually the reaction in each of the reactors.
This invention is applied most effectively to such a
10 case because it enables to know and control individually
the respective polymer composition in each of the reactors,
and thus to produce the objective polymer stably.

This invention is applied effectively to a case
where polymerization is conducted by continuous operation,
15 but it may also be applied to non-stationary operations
such as batch-wise polymerization. The application of
the technique of this invention enables, even in
batch-wise polymerization, to know the composition and
the amount of polymer formed at time intervals during the
20 progress of polymerization from material balance,
heat balance, and monomer reactivity ratio, and thus to
control the polymerization stably.

Thus, this invention is in no way restricted
with respect to methods of polymerization, polymerization
25 conditoinis, and forms of operation, etc. The invention is
further described in detail below with reference to the
following Example.

1 Example

An ethylene-butene-1 copolymer was produced by a continuous operation using the polymerization equipment shown in the attached drawing.

5 In a stationary state, an organoaluminum catalyst, a transition metal compound catalyst containing titanium, hydrogen, ethylene, butene-1 and butane were fed at constant rates to the reactor through the line 5. The temperatures, pressures, and flow rates of the raw
10 materials being fed were measured separately. The polymerization was continued withdrawing a corresponding amount of the reaction mixture to those of the fed materials. The reactor was operated in the substantial absence of a gas phase.

15 The temperature of the reactor 1 was controlled by the cooling water circulated by the pump 3 to keep a constant value. The temperature of the circulating water was controlled by introducing a cooling water of lower temperature from the line 7 and withdrawing a corres-
20 ponding amount of a cooling water of higher temperature through the line 8. The physical quantities measured in each of the lines and instruments were input to the computer 4 to be subjected to the calculation of the material and heat balance.

25 Here, with respect to ethylene and butene-1 which participate in the reaction of this invention, the fed amount, the heat of reaction, the reactivity ratio and other quantities are defined as follows:

	Fed amount (line 5) kg/H	Unreacted amount (line 6) kg/H	Polymerized amount (line 6) kg/H	Heat of polymerization kcal/kg
Ethylene	F_1	a	A	h_1
Butene-1	F_2	b	B	h_2

1 The reactivity ratio $(A/B)/(a/b) = r$.

Here, F_1 and F_2 are measured values, and h_1 , h_2 , and r are values obtained beforehand.

Measured amount of heat: Q kcal/H.

5 Here, Q represents the amount of heat evolved by polymerization alone, which can be obtained by subtracting, from the total amount of heat calculated from the heat balance of cooling water, the amounts of such heat other than that due to the polymerization as
10 the sensible heat of feed materials, the heat required to keep the reaction mixture in the reactor flowing, and the heat dissipated from the reactor.

From the material balance:

$$F_1 = a + A \quad (1)$$

15 $F_2 = b + B \quad (2)$

From the reactivity ratio:

$$(A/B)/(a/b) = r \quad (3)$$

From the heat balance:

$$Q = h_1 A + h_2 B \quad (4)$$

20 From equations (1) to (4), the following equations can be set up:



$$A = \frac{-h_1 F_1 r - h_2 F_2 + Q(1-r) + \sqrt{\{h_1 F_1 r + h_2 F_2 - Q(1-r)\}^2 + 4h_1(1-r)QF_1 r}}{2h_1(1-r)} \quad (5)$$

$$B = F_2 A / \{(F_1 - A)r + A\} \quad (6)$$

1 The heats of polymerization of ethylene and
butene-1 are:

$$h_1 = 910 \text{ kcal/kg}, \quad h_2 = 360 \text{ kcal/kg}.$$

5 A separate experiment under the same conditions
as those in this Example gave:

$$r = 50.$$

At a certain point of time in the polymerization
of, this Example, the flow rate in the line 5 and Q had
the following values:

$$\begin{aligned} 10 \quad F_1 &= 10 \text{ kg/H} \\ F_2 &= 3.7 \text{ kg/H} \\ Q &= 8624 \text{ kcal/H} \end{aligned}$$

From equations (5) and (6), by the use of the
computer 4, A and B were obtained as follows:

$$15 \quad A = 9.2 \text{ kg/H}, \quad B = 0.7 \text{ kg/H}.$$

Consequently, the butene-1 content of the polymer and
the polymerized amount were calculated at 7.1% by weight
and 9.9 kg/H, respectively.

1 On the other hand, the objective values in
this Example were as follows:

Butene-1 content of the polymer, 8.0% by weight:

Polymerized amount, 10 kg/H.

5 The objective values can be attained, while keeping the
amount of feed ethylene at 10 kg/H, by giving Q and E
the following values:

$$Q = 8660 \text{ kcal/H}$$

$$F_2 = 4.3 \text{ kg/H}$$

10. Accordingly, the amount of butene-1 being fed was increased
a little, and further the amount of the catalyst being
fed was increased so as to make Q 8660 kcal/H. These
operations were controlled automatically by the outputs
from the computer.

15 Owing to the above alteration of reaction condi-
tions, the difference between the temperature inside the
reactor and that of the jacket, ΔT , increased a little.
The change of ΔT was monitored to confirm that the poly-
merization was proceeding as intended. Further, the mate-
20 rial balance was determined from the flow rates and
difference of temperatures in lines 7 and 8, to confirm
the progress of polymerization thereby.

After a while, the intended polymer has come to
be obtained stably.

WHAT IS CLAIMED IS:

1. In a process for producing an intended polymer by polymerizing at least one vinyl compound, an improvement which comprises calculating the concentration of the unreacted vinyl compound in the reactor, the composition of the polymer formed, and the amount thereof from the amount of heat calculated from the heat balance of the reactor, the amount of raw materials fed to the reactor, the reactivity ratio of the vinyl compounds and the heat of polymerization per unit amount of each of the vinyl compounds, and regulating the amounts of raw materials to be fed into the reactor in such a way that the values thus calculated agree respectively with the predetermined target values of the polymer composition and the amount of polymer formed.
2. A process according to Claim 1, wherein said calculation and regulation are performed constantly by using a computer.
3. A process according to Claim 1, wherein the difference between the temperature inside the reactor and that of the reactor jacket is used as a parameter in the calculation of the amount of reaction heat.
4. A process according to Claim 1, wherein the flow rate of a cooling medium supplied into the reaction jacket from outside of the system as well as the difference in temperatures of said cooling medium at the inlet and at the outlet are used as parameters in the calculation of the amount of reaction heat.

5. A process according to Claim 1, wherein a copolymerization is carried out by using plural vinyl compounds to be polymerized.
6. A process according to Claim 1, wherein the vinyl compound is a 1-olefin.
7. A process according to Claim 6, wherein the 1-olefin is ethylene and/or a 1-olefin having 3 to 12 carbon atoms and a Ziegler-Natta catalyst is used as the catalyst.
8. A process according to Claim 1, wherein the reactor is one having plural reactors connected in series.
9. A process according to Claim 1, wherein the polymerization is a solution or slurry polymerization.
10. A process according to Claim 1, wherein the polymerization is a gas-phase polymerization.
11. A process according to Claim 1, wherein the reactor is operated in a state filled with a liquid.